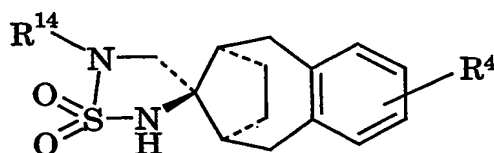


CLAIMS

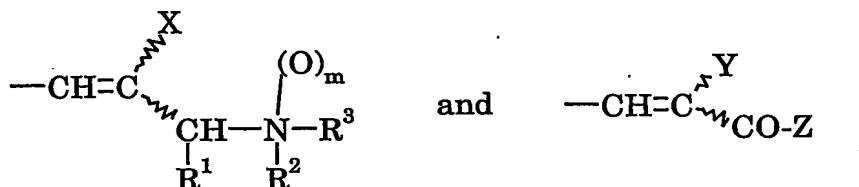
1. A compound of formula I:



I

5

wherein R⁴ is selected from:



X represents H, halogen, CN or methyl;

R¹ represents H or C₁₋₄alkyl which is optionally substituted with OH or C₁₋₄alkoxy; or R¹ and R² together complete a heterocyclic ring of 3-7 members bearing 0-2 substituents, in addition to R³, selected from halogen, oxo, NO₂, CN, CF₃, C₁₋₆alkyl, C₂₋₆acyl, C₂₋₆alkenyl, C₁₋₆alkoxy, C₁₋₆alkoxycarbonyl and Ar;

when R¹ represents H or optionally substituted C₁₋₄alkyl, R² and R³ independently represent H, C₁₋₁₀alkyl, C₃₋₁₀cycloalkyl, C₃₋₆cycloalkylC₁₋₆alkyl, C₂₋₁₀alkenyl, C₂₋₁₀alkynyl, Ar, heterocyclyl, or heterocyclylC₁₋₆alkyl, wherein the alkyl, cycloalkyl, alkenyl and alkynyl groups optionally bear one substituent selected from halogen, CF₃, NO₂, CN, Ar, ArCH₂O, ArO, -OR¹¹, -SR¹¹, -SO₂R¹², -COR¹¹, -CO₂R¹¹, -CON(R¹¹)₂, -OCOR¹², -N(R¹¹)₂ and -NR¹¹COR¹²; and the heterocyclic groups optionally bear up to 3 substituents independently selected from halogen, NO₂, CN, R¹², Ar, ArCH₂O, ArO, ArOCH₂, -OR¹¹, -SR¹¹, -SO₂R¹², -COR¹¹, -CO₂R¹¹, -CON(R¹¹)₂, -OCOR¹², -N(R¹¹)₂ and -NR¹¹COR¹²;

or R² and R³ together with the nitrogen to which they are mutually attached complete a mono- or bicyclic heterocyclic ring system of 5-10 ring

atoms selected from C, N, O and S, said ring system optionally having an additional benzene or heteroaryl ring fused thereto, said heterocyclic system and optional fused ring bearing 0-3 substituents independently selected from halogen, oxo, NO₂, CN, R¹², Ar, ArCH₂O, ArO, ArOCH₂,
5 -OR¹¹, -SR¹¹, -SO₂R¹², -COR¹¹, -CO₂R¹¹, -CON(R¹¹)₂, -OCOR¹², -N(R¹¹)₂ and -NR¹¹COR¹²;

and when R¹ completes a ring with R², R³ represents H, C₁₋₆alkyl, C₂₋₆acyl, C₂₋₆alkenyl or benzyl;

m is 0 or 1, with the proviso that when m is 1 neither R² nor R³ is H
10 and R³ is not acyl, and that m is 1 when X and R¹ are both H;

R¹¹ represents H or R¹²;

R¹² represents C₁₋₆alkyl which optionally bears up to 3 halogen substituents or one substituent selected from CN, OH, C₁₋₄alkoxy and C₁₋₄alkoxycarbonyl;

15 Y represents halogen, CN or methyl;

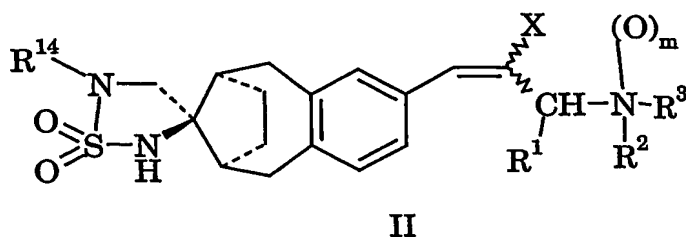
Z represents OR¹¹ or N(R⁵)R⁶;

R⁵ and R⁶ have the same definition as R² and R³ in the embodiment in which R¹ is H or optionally substituted C₁₋₄alkyl;

R¹⁴ represents H or C₁₋₆alkyl, C₃₋₇cycloalkyl, C₃₋₆cycloalkylC₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, phenyl or benzyl, any of which optionally bear up to 3 halogen substituents or one substituent selected from CN, NO₂, OH, C₁₋₄alkoxy, CO₂H, C₁₋₄alkoxycarbonyl, C₂₋₆acyl, C₂₋₆acyloxy, amino, C₁₋₄alkylamino, di(C₁₋₄alkyl)amino, C₂₋₆acylamino, carbamoyl, C₁₋₄alkylcarbamoyl and di(C₁₋₄alkyl)carbamoyl; and
20

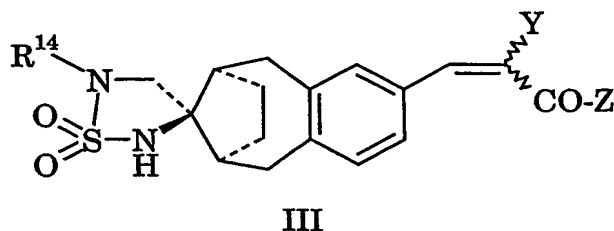
25 Ar represents phenyl or heteroaryl either of which optionally bears up to 3 substituents independently selected from halogen, CF₃, NO₂, CN, OCF₃, C₁₋₆alkyl and C₁₋₆alkoxy;
or a pharmaceutically acceptable salt thereof.

30 2. A compound according to claim 1 of formula II:



or a pharmaceutically acceptable salt thereof.

- 5 3. A compound according to claim 2 wherein R¹ and R² complete a heterocyclic ring of 5 or 6 atoms and R³ represents H, C₁₋₆alkyl, C₂₋₆acyl or benzyl.
4. A compound according to claim 2 wherein R¹ is H or optionally substituted C₁₋₄alkyl and R² and R³ complete a heterocyclic ring system.
- 10 5. A compound according to claim 4 wherein R¹⁴ is 2,2,2-trifluoroethyl, X is F, CN or methyl, and R¹ is H.
- 15 6. A compound according to claim 4 wherein m is 1 and X and R¹ are both H.
7. A compound according to claim 1 of formula III:



or a pharmaceutically acceptable salt thereof.

8. A compound according to claim 7 wherein Y represents F, CN or methyl and Z represents OH, C₁₋₆alkoxy or N(R⁵)R⁶.
9. A compound according to claim 8 wherein R¹⁴ represents 2,2,2-trifluoroethyl and Z represents ethoxy.
10. A pharmaceutical composition comprising a compound according to any previous claim and a pharmaceutical carrier.
11. A compound according to any of claims 1-9 for use in a method of treatment of the human body.
12. The use of a compound according to any of claims 1-9 in the manufacture of a medicament for treating or preventing Alzheimer's disease.
13. A method of treatment of a subject suffering from or prone to Alzheimer's disease comprising administering to that subject an effective amount of a compound according to any of claims 1-9.